

**Amendments to the Claims:**

The following claims will replace all prior versions of the claims in this application (in the unlikely event that no claims follow herein, the previously pending claims will remain):

1. (Currently amended) A method for the production of  $\alpha$ -L-aspartyl-L-phenylalanine (Asp-Phe) from the substrates L-aspartic acid (L-Asp) and L-phenylalanine (L-Phe) which comprises (a) contacting the substrates in the presence of an effective amount of adenosine-triphosphate (ATP), with a non-ribosomal dipeptide synthetase comprising two minimal modules connected by one condensation domain, wherein the N-terminal module of these minimal modules recognizes L-aspartic acid, wherein ~~and~~ the C-terminal module of these minimal modules recognizes L-phenylalanine and is covalently bound at its N-terminal end to the condensation domain, and wherein each of these minimal modules ~~is composed of~~ comprises an adenylation domain and a thiolation domain containing a 4'-phosphopantetheinyl cofactor, wherein the dipeptide synthetase further comprises a thioesterase releasing factor for facilitating release of the  $\alpha$ -L-aspartyl-L-phenylalanine (Asp-Phe) dipeptide, and wherein the adenylation domains are identifiable by having at least substantial structural homology with the core motifs having SEQ ID NOs: 1, 2, 3, 4, 5, 6, 7, 8, 9 and 10 found in naturally occurring non-ribosomal peptide synthetases and (b) recovering the  $\alpha$ -L-aspartyl-L-phenylalanine (Asp-Phe) produced in (a).
2. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 1, wherein the condensation domain in the dipeptide synthetase is also covalently bound to the module recognising L-aspartic acid.
3. (Cancelled).
4. (Currently Amended) ~~Method~~ The method for the production of Asp-Phe according to claim ~~3~~ 1, wherein the condensation domain in the dipeptide synthetase is covalently bound to the module recognizing L-aspartic acid and wherein the thioesterase releasing factor is covalently bound to the module recognizing L-phenylalanine.
5. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 1, wherein a non-integrated protein with thioesterase Type-II activity is further present together with the dipeptide synthetase.

6. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 5, wherein the dipeptide synthetase is present in a microorganism, said process further comprising growing said microorganism in a fermentor and feeding glucose, L-Asp, L-Phe, or mixtures thereof to said fermentor.

7. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 6, wherein the microorganism is first grown in a fermentor to reach a predetermined cell density before the expression of the Asp-Phe dipeptide synthetase is ~~switched-on~~ induced, and wherein the glucose, L-Asp, L-Phe, or mixture thereof is added at the same time the expression of the Asp-Phe dipeptide is ~~switched-on~~ induced.

8. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 7, wherein the microorganism is an L-phenylalanine producing microorganism, and only the glucose and L-Asp are fed.

9. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 8, wherein the microorganism is an *Escherichia* or *Bacillus* species.

10. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 6, wherein the microorganism used is a strain having reduced protease activity for Asp-Phe or having no protease activity towards Asp-Phe.

11. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 1, wherein the production of Asp-Phe is carried out using the dipeptide synthetase in its isolated form in a reactor and simultaneously supplying L-Asp, L-Phe, or a mixture thereof and ATP to the reactor.

12. (Currently presented) ~~Method~~ The method for the production of Asp-Phe according to claim 11, wherein the supply of ATP is provided in part by an in situ ATP-regenerating system.

13. (Currently amended) ~~Method~~ The method for the production of Asp-Phe according to claim 12, wherein the ATP-regenerating system is present in a permeabilised microorganism.

14. (Withdrawn) A DNA fragment or a combination of DNA fragments coding for a non-ribosomal Asp-Phe dipeptide synthetase, said synthetase comprising two minimal modules connected by one condensation domain, wherein the N-terminal module of these minimal modules recognizes L-aspartic acid, and the C-terminal module of these minimal modules recognizes L-phenylalanine and is covalently bound at its N-terminal end to the condensation domain, and wherein each of said minimal modules is composed of an adenylation domain and a thiolation domain containing a 4'-phosphopantetheinyl cofactor.

15. (Withdrawn) A DNA fragment coding for an Asp-Phe dipeptide synthetase according to claim 14, wherein the condensation domain in the encoded dipeptide synthetase is also covalently bound to the module recognising L-aspartic acid.

16. (Withdrawn) A DNA fragment or a combination of DNA fragments according to claim 14, wherein the DNA fragment or the combination of DNA fragments encoding the dipeptide synthetase also code for a thioesterase releasing factor for the Asp-Phe formed on that dipeptide synthetase.

17. (Withdrawn) A DNA fragment according to claim 16, wherein the condensation domain in the dipeptide synthetase is covalently bound to the module recognizing L-aspartic acid and wherein the thioesterase releasing factor is covalently bound to the module recognizing L-phenylalanine.

18. (Withdrawn) A DNA fragment or a combination of DNA fragments according to claim 14, wherein said DNA fragment or a combination of DNA fragments also code for a non-integrated protein with thioesterase Type-II activity.

19. (Withdrawn) A recombinant microorganism containing a DNA fragment or a combination of DNA fragments according to claim 14.

20. (Withdrawn) A microorganism according to claim 19, wherein the microorganism is capable of producing L-Asp, L-Phe, or a mixture thereof.

21. (Withdrawn) A micro-organism according to claim 20, wherein the micro-organism is an *Escherichia coli* or *Bacillus* species.

22. (Withdrawn) Non-ribosomal Asp-Phe dipeptide synthetase comprising two minimal modules connected by one condensation domain, wherein the N-terminal module of these minimal modules recognizes L-aspartic acid and the C-terminal module of these minimal modules recognizes L-phenylalanine and is covalently bound at its N-terminal end to the condensation domain, and wherein each of these minimal modules is composed of an adenylation domain and a thiolation domain containing a 4'-phosphopantetheinyl cofactor.

23. (Withdrawn) Non-ribosomal Asp-Phe dipeptide synthetase according to claim 22, wherein the condensation domain in the dipeptide synthetase is also covalently bound to the module recognizing L-aspartic acid.

24. (Withdrawn) Non-ribosomal Asp-Phe dipeptide synthetase according to claim 22, wherein the dipeptide synthetase also comprises a releasing factor for the Asp-Phe formed on that dipeptide synthetase.

25. (Withdrawn) Non-ribosomal Asp-Phe dipeptide synthetase according to claim 24, wherein the condensation domain in the dipeptide synthetase is covalently bound to the module recognizing L-aspartic acid and wherein the releasing factor is covalently bound to the module recognizing L-phenylalanine.

26. (Currently amended) A method for the production of  $\alpha$ -L-aspartyl-L-phenylalanine (Asp-Phe) from the substrates L-aspartic acid (L-Asp) and L-phenylalanine (L-Phe) which comprises (a) contacting the substrates, in the presence of an effective amount of adenosine-triphosphate (ATP), with a non-ribosomal dipeptide synthetase comprising two minimal modules, one minimal module being encoded by DNA comprising part of the *srfB* gene from *B. subtilis* ATCC 21332 recognizing L-aspartic acid and the second minimal module being encoded by DNA comprising part of the *tycA* gene from *B. brevis* ATCC 8185 recognizing L-phenylalanine, the two minimal modules being connected by one condensation domain, wherein the N-terminal module of these minimal modules recognizes L-aspartic acid, wherein and the C-terminal module of these minimal modules recognizes L-phenylalanine and is covalently bound at its N-terminal end to the condensation domain, and wherein each of these minimal modules ~~is composed of~~ comprises an adenylation domain and a thiolation domain containing a 4'-phosphopantetheinyl cofactor, wherein the dipeptide synthetase further comprises a thioesterase releasing factor for facilitating release of the  $\alpha$ -

L-aspartyl-L-phenylalanine (Asp-Phe) dipeptide, and wherein the adenylation domains are identifiable by having at least substantial structural homology with the core motifs having SEQ ID NOs: 1, 2, 3, 4, 5, 6, 7, 8, 9 and 10 found in naturally occurring non-ribosomal peptide synthetases, and (b) recovering the  $\alpha$ -L-aspartyl-L-phenylalanine (Asp-Phe) produced in (a).